=> file reg FILE 'REGISTRY' ENTERED AT 20:31:26 ON 15 FEB 2006 USE IS SUBJECT TO THE TERMS OF YOUR STN CUSTOMER AGREEMENT. PLEASE SEE "HELP USAGETERMS" FOR DETAILS. COPYRIGHT (C) 2006 American Chemical Society (ACS) => d his FILE 'LREGISTRY' ENTERED AT 13:50:33 ON 15 FEB 2006 L1STR FILE 'HCAPLUS' ENTERED AT 14:09:17 ON 15 FEB 2006 380721 S WANG ?/AU L2L3 19509 S MAO ?/AU L42489 S LEONG ?/AU 27 S L2 AND L3 AND L4 L5 L6 46365 S WANG J?/AU L7 1615 S MAO H?/AU 548 S LEONG K?/AU L8 L9 19 S L6 AND L7 AND L8 SEL L9 1,2,4,6,7,11,12,13,14,15,16,18,19 RN FILE 'REGISTRY' ENTERED AT 14:19:24 ON 15 FEB 2006 40 S E1-E40 L10L1110 S L10 AND P/ELS L128 S L10 AND PMS/CI L13 7 S L11 AND L12 FILE 'HCAPLUS' ENTERED AT 14:21:06 ON 15 FEB 2006 SEL L9 1-19 RN FILE 'REGISTRY' ENTERED AT 14:21:27 ON 15 FEB 2006 L1453 S E41-E93 L15 12 S L14 AND P/ELS L16 14 S L14 AND PMS/CI L17 8 S L15 AND L16 L18 8 S L17 OR L13 L19 30 S L14 AND N/ELS L20 3 S L19 AND L15 FILE 'HCAPLUS' ENTERED AT 15:09:28 ON 15 FEB 2006

FILE 'REGISTRY' ENTERED AT 15:13:22 ON 15 FEB 2006 1 S 168788-11-2 1 S 100-36-7

24 S L18

13 S L9 AND L21

L21 L22

L23

L24

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L25
             1 S 107-15-3
L26
             1 S 108-00-9
L27
             1 S 109-81-9
L28
             1 S 51460-23-2
             1 S 18807-71-1
L29
L30
             1 S 162576-01-4
L31
             1 S 110-60-1
             1 S 15302-88-2
L32
L33
             1 S 109-76-2
             1 S 38170-37-5
L34
             1 S 77987-49-6
L35
L36
             1 S 17996-12-2
L37
             1 S 622-08-2
L38
             1 S 158474-89-6
L39
             16 S L23-L38
     FILE 'HCA' ENTERED AT 16:41:57 ON 15 FEB 2006
          45254 S L39
L40
L41
             10 S L21 AND L40
     FILE 'LREGISTRY' ENTERED AT 16:42:52 ON 15 FEB 2006
L42
                STR
     FILE 'REGISTRY' ENTERED AT 16:58:11 ON 15 FEB 2006
L43
               SCR 2043
L44
            50 S L42 AND L43
     FILE 'LREGISTRY' ENTERED AT 16:59:13 ON 15 FEB 2006
L45
                STR L42
     FILE 'REGISTRY' ENTERED AT 17:19:20 ON 15 FEB 2006
L46
             0 S L45 AND L43
     FILE 'LREGISTRY' ENTERED AT 17:19:57 ON 15 FEB 2006
L47
               STR L45
     FILE 'REGISTRY' ENTERED AT 17:22:10 ON 15 FEB 2006
            50 S L47 AND L43
L48
L49
           4964 S L47 AND L43 FUL
               SAV L49 ACO898/A
L50
             50 S L42 SSS SAM SUB=L49
L51
          1052 S L42 SSS FUL SUB=L49
               SAV L51 ACQ898A/A
    FILE 'HCA' ENTERED AT 18:57:01 ON 15 FEB 2006
L52
           536 S L51
L53
           111 S L51/D OR L51/DP
L54
          7267 S L39/D OR L39/DP
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FILE 'REGISTRY' ENTERED AT 18:58:08 ON 15 FEB 2006
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L56
             3 S L39 AND ?HYDROXY?/CNS
L57
             13 S L39 NOT L56
     FILE 'HCA' ENTERED AT 19:00:57 ON 15 FEB 2006
L58
          44419 S L57
           7258 S L57/D OR L57/DP
L59
             9 S L52 AND L58
L60
L61
             5 S L53 AND L59
           866 S L56
L62
L63
             9 S L56/D OR L56/DP
             5 S L52 AND L62
L64
             4 S L53 AND L63
L65
L66
            13 S L53 (L) (AMINAT? OR POLYAMINAT? OR AMINE# OR AMINO# OR
            7 S L53 (L) (?ETHYLENEDIAMIN? OR ?AMINOMETHYL? OR ?AMINOETH
L67
L68
            19 S L52 (L) (AMINAT? OR POLYAMINAT? OR AMINE# OR AMINO# OR
            10 S L52 (L) (?ETHYLENEDIAMIN? OR ?AMINOMETHYL? OR ?AMINOETH
L69
L70
           311 S ?BENZYLOXYETHANOL?
              1 S L52 AND L70
L71
     FILE 'REGISTRY' ENTERED AT 19:12:16 ON 15 FEB 2006
L72
          36080 S (C(L)H(L)N)/ELS (L) 3/ELC.SUB AND NO RSD/FA
L73
          20535 S L72 AND 1<N
     FILE 'HCA' ENTERED AT 19:13:20 ON 15 FEB 2006
        172109 S L73
L74
         27070 S L73/D OR L73/DP
L75
L76
            19 S L52 AND L74
L77
             8 S L53 AND L75
     FILE 'REGISTRY' ENTERED AT 19:15:51 ON 15 FEB 2006
        374401 S (C(L)H(L)N(L)O)/ELS (L) 4/ELC.SUB AND NO RSD/FA
L78
L79
        1680139 S (C(L)H(L)N(L)O)/ELS (L) 4/ELC.SUB AND 1/NRS
L80
         54912 S L79 AND ?CARBAM?/CNS
L81
         42533 S L80 AND 2<0
        374401 S L78 OR L78
L82
L83
        194401 S L82 RAN=(,159430-98-5)
L84
        180000 S L82 RAN=(159431-21-7,)
     FILE 'HCA' ENTERED AT 19:34:05 ON 15 FEB 2006
L85
          1916 S L81/D OR L81/DP
L86
         85768 S L83/D OR L83/DP
L87
          3492 S L84/D OR L84/DP
L88
            17 S L53 AND (L85 OR L86 OR L87)
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FILE 'LREGISTRY' ENTERED AT 19:38:03 ON 15 FEB 2006

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L89
               STR L47
    FILE 'REGISTRY' ENTERED AT 19:40:09 ON 15 FEB 2006
L90
             4 S L89 SSS SAM SUB=L49
L91
           143 S L89 SSS FUL SUB=L49
               SAV L91 ACQ565/A
    FILE 'HCA' ENTERED AT 20:06:37 ON 15 FEB 2006
           44 S L91
L92
            29 S L41 OR L60 OR L61 OR L66 OR L67 OR L77
L93
L94
           57 S (L68 OR L69 OR L76 OR L92) NOT L93
L95
           19 S L93 AND (1840-2001/PRY OR 1840-2001/PY)
L96
           47 S L94 AND (1840-2001/PRY OR 1840-2001/PY)
           10 S L93 NOT L95
L97
               SAV L97 ACQ565A/A
L98 10 S L94 NOT L96
               SAV L98 ACQ565B/A
    FILE 'LREGISTRY' ENTERED AT 20:12:19 ON 15 FEB 2006
L99
               STR L47
    FILE 'REGISTRY' ENTERED AT 20:18:23 ON 15 FEB 2006
           39 S L99 SSS SAM SUB=L49
L100
L101
               STR
          6 S (L99 NOT L101) SSS SAM SUB=L49
L102
L103
          148 S (L99 NOT L101) SSS FUL SUB=L49
               SAV L103 ACQ898B/A
    FILE 'HCA' ENTERED AT 20:25:22 ON 15 FEB 2006
L104
           83 S L103
L105
            19 S L64 OR L65 OR L71 OR L88
L106
           83 S L104 NOT L105
L107
           14 S L105 AND (1840-2001/PRY OR 1840-2001/PY)
            74 S L106 AND (1840-2001/PRY OR 1840-2001/PY)
L108
L109
            5 S L105 NOT L107
               SAV L109 ACQ898C/A
             9 S L106 NOT L108
L110
               SAV L110 ACQ898D/A
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FILE 'REGISTRY' ENTERED AT 20:31:26 ON 15 FEB 2006

=> d 191 que stat L43 SCR 2043 L47 STR

Page 1-A

Page 1-B

VAR G1=C/ID

VAR G2=7/12/16/20/25/29

REP G3 = (1-6) CH2

NODE ATTRIBUTES:

HCOUNT IS E1 AT 12
HCOUNT IS E1 AT 16
NSPEC IS RC AT 32
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 20

STEREO ATTRIBUTES: NONE

L49 4964 SEA FILE=REGISTRY SSS FUL L47 AND L43 L89 STR

21 0 || 1 P-X-0X G1 2 3 N 33

VAR G1=C/ID NODE ATTRIBUTES: DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 5

STEREO ATTRIBUTES: NONE

L91 143 SEA FILE=REGISTRY SUB=L49 SSS FUL L89

100.0% PROCESSED 157 ITERATIONS 143 ANSWERS

SEARCH TIME: 00.00.01

=> file hca
FILE 'HCA' ENTERED AT 20:32:01 ON 15 FEB 2006
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ANSWER 1 OF 19 HCA COPYRIGHT 2006 ACS on STN

Society: Minneapolis, Minn. (English) 2001.

=> d 195 1-19 cbib abs hitstr hitind

138:112274 Biodegradable polyphosphoramidates as gene carriers: Effect of charge groups on transfection efficiency. Wang, J.; Mao, H.-Q.; Leong, K. W. (Clinical Research Center, Johns Hopkins Singapore, Singapore, 117597, Singapore). Proceedings - 28th International Symposium on Controlled Release of Bioactive Materials and 4th Consumer & Diversified Products Conference, San Diego, CA, United States, June 23-27, 2001, Volume 2, 1119-1120. Controlled Release

CODEN:

- AB To elucidate the important features of polymeric gene carriers, a series of new cationic polyphosphoramidates (PPAs) with an identical backbone but different side chains contg. primary to tertiary amino groups were synthesized. These PPA carriers showed different transfection abilities and DNA binding capacities. PPA with a spermidine side chain was the most efficient in several cell lines, while other PPAs only achieved moderate levels of gene expression. Chloroquine significantly enhanced the transfection efficiency. These PPAs were less toxic than PEI and PLL.
- 100-36-7DP, reaction products with polydioxaphospholane 108-00-9DP, reaction products with polydioxaphospholane 109-76-2DP, 1,3-Propanediamine, reaction products with polydioxaphospholane 38170-37-5DP, reaction products with polydioxaphospholane 51460-23-2DP, reaction products with polydioxaphospholane 83945-68-0DP, reaction products with amines

(charge groups effect on transfection efficiency of biodegradable polyphosphoramidates as gene carriers)

RN 100-36-7 HCA

CN 1,2-Ethanediamine, N,N-diethyl- (9CI) (CA INDEX NAME)

Et₂N-CH₂-CH₂-NH₂

RN 108-00-9 HCA

CN 1,2-Ethanediamine, N,N-dimethyl- (9CI) (CA INDEX NAME)

 $Me_2N-CH_2-CH_2-NH_2$

RN 109-76-2 HCA

CN 1,3-Propanediamine (6CI, 8CI, 9CI) (CA INDEX NAME)

 $H_2N-CH_2-CH_2-CH_2-NH_2$

RN 38170-37-5 HCA

CN Ethanaminium, 2-amino-N,N,N-trimethyl- (9CI) (CA INDEX NAME)

 $Me_3+N-CH_2-CH_2-NH_2$

RN 51460-23-2 HCA

CN 1,4-Butanediamine, N-(3-aminopropyl)-N-methyl- (9CI) (CA INDEX NAME)

 $^{\rm Me}$ $^{\rm |}$ $^{\rm H}_{\rm 2}{\rm N}^{-}$ (CH₂) $^{\rm 3}_{\rm 3}^{-}$ N $^{\rm -}$ (CH₂) $^{\rm 4}_{\rm 4}^{-}$ NH₂

RN 83945-68-0 HCA

CN 1,3,2-Dioxaphospholane, 4-methyl-, 2-oxide, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 16352-26-4 CMF C3 H7 O3 P

O H O Me

- CC 63-6 (Pharmaceuticals)
 Section cross-reference(s): 3
- 100-36-7DP, reaction products with polydioxaphospholane 108-00-9DP, reaction products with polydioxaphospholane 109-76-2DP, 1,3-Propanediamine, reaction products with polydioxaphospholane 38170-37-5DP, reaction products with polydioxaphospholane 51460-23-2DP, reaction products with polydioxaphospholane 83945-68-0DP, reaction products with amines

(charge groups effect on transfection efficiency of biodegradable polyphosphoramidates as gene carriers)

- L95 ANSWER 2 OF 19 HCA COPYRIGHT 2006 ACS on STN
- 138:112127 Prolonged and enhanced gene expression in mouse muscle by sustained release of plasmid DNA using PPE as a carrier. Wang, J.; Mao, H.-Q.; Leong, K. W. (Johns Hopkins Singapore, Singapore, 117597, Singapore). Proceedings 28th International Symposium on Controlled Release of Bioactive Materials and 4th Consumer & Diversified Products Conference, San Diego, CA, United States, June 23-27, 2001, Volume 1, 25-26. Controlled Release Society: Minneapolis, Minn. (English) 2001. CODEN: 69CNY8.
- AB A new system for the controlled release of plasmid DNA was developed using a biodegradable polyphosphoester [PPE]. The release of plasmid DNA from PPE-DNA complexes was a function of charge ratio. A an N/P ratio of 1, a burst of 14.1 .mu.g/mL was obsd. for the first day, followed by a near const. release of 1.24 .mu.g/mL/day for 12 days. Higher charge ratios reduced both the DNA release rate and the burst effect. A 20 and 16-fold higher levels of .beta.-Gal expression in mouse muscle were achieved at day 7 using PPE-DNA complexes with N/P ratios of 0.5 and 1 as compared with naked DNA injection, and the gene expression level was maintained for up to 4 wk.
- IT 31259-60-6P

(prolonged and enhanced gene expression in mouse muscle by sustained release of plasmid DNA using polyphosphoester as a carrier)

- RN 31259-60-6 HCA
- CN 1,3,2-Dioxaphospholane, 2-chloro-4-methyl-, 2-oxide, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 4225-53-0 CMF C3 H6 Cl O3 P

T77987-49-6, Benzyl N-(2-hydroxyethyl)carbamate (prolonged and enhanced gene expression in mouse muscle by sustained release of plasmid DNA using polyphosphoester as a carrier)

RN 77987-49-6 HCA

CN Carbamic acid, (2-hydroxyethyl)-, phenylmethyl ester (9CI) (CA INDEX NAME)

RN 31259-60-6 HCA

CN 1,3,2-Dioxaphospholane, 2-chloro-4-methyl-, 2-oxide, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 4225-53-0 CMF C3 H6 Cl O3 P

RN 77987-49-6 HCA

CN Carbamic acid, (2-hydroxyethyl)-, phenylmethyl ester (9CI) (CA INDEX NAME)

- CC 63-5 (Pharmaceuticals)
- IT 31259-60-6P

(prolonged and enhanced gene expression in mouse muscle by sustained release of plasmid DNA using polyphosphoester as a carrier)

- T77987-49-6, Benzyl N-(2-hydroxyethyl)carbamate (prolonged and enhanced gene expression in mouse muscle by sustained release of plasmid DNA using polyphosphoester as a carrier)
- IT 31259-60-6DP, aminoethoxy derivs.
 - 77987-49-6DP, reaction products with polydioxaphospholane (prolonged and enhanced gene expression in mouse muscle by sustained release of plasmid DNA using polyphosphoester as a carrier)
- L95 ANSWER 3 OF 19 HCA COPYRIGHT 2006 ACS on STN
- 138:78460 Biodegradable polyphosphoramidates for controlled release of bioactive substances. Wang, Jun; Mao, Hai-Quan; Leong, Kam Weng (Johns Hopkins Singapore Pte. Ltd., Singapore). PCT Int. Appl. WO 2003000776 Al 20030103, 52 pp. DESIGNATED STATES: W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: PIXXD2. APPLICATION: WO 2002-SG91 20020514. PRIORITY: US 2001-PV290833 20010514.
- The present invention is directed to a series of new polycationic biodegradable polyphosphoramidates. A process for making the polymers, compns. contg. these polymers and bioactive ligands to enhance the cellular uptake ad intracellular trafficking, articles and methods for delivery of drugs and genes using these polymers are described. A gene delivery system based on these polymers is prepd. by complex coacervation of nucleic acid (DNA or RNA) with polymers. Targeting ligands and mols. that could facilitate gene transfer can be conjugated to polymers to achieve selective and enhanced gene delivery. The current invention also provides a complex compn. with buffering capacity. [OCHMeCH2OP(O)H]n is prepd. and treated with a no. of polyamines to form the polyphosphoramidates. Examples also

give include cytotoxicity assay, gel retardation assay for the DNA binding capacity, and prepn. of DNA and chloroquine sulfate coacervates with the polyphosphoramidates.

IT 83945-68-0P

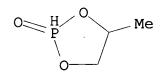
(biodegradable polyphosphoramidates for controlled release of bioactive substances)

RN 83945-68-0 HCA

CN 1,3,2-Dioxaphospholane, 4-methyl-, 2-oxide, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 16352-26-4 CMF C3 H7 O3 P



IT 100-36-7DP, N,N-Diethylethylenediamine, reaction products
with poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
108-00-9DP, N,N-Dimethylethylenediamine, reaction products
with poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
110-60-1DP, 1,4-Butanediamine, reaction products with
poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
15302-88-2DP, reaction products with poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
120527-66-4DP, reaction
products with polyamines 168788-11-2DP, reaction
products with poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
(biodegradable polyphosphoramidates for controlled release of bioactive substances)

RN 100-36-7 HCA

CN 1,2-Ethanediamine, N,N-diethyl- (9CI) (CA INDEX NAME)

Et₂N-CH₂-CH₂-NH₂

RN 108-00-9 HCA

CN 1,2-Ethanediamine, N,N-dimethyl- (9CI) (CA INDEX NAME)

 $Me_2N-CH_2-CH_2-NH_2$

RN 110-60-1 HCA

CN 1,4-Butanediamine (8CI, 9CI) (CA INDEX NAME)

 $H_2N-(CH_2)_4-NH_2$

RN 15302-88-2 HCA

CN Ethanaminium, N,N,N-trimethyl- (9CI) (CA INDEX NAME)

RN 120527-66-4 HCA

CN Poly[oxyphosphinylideneoxy(methyl-1,2-ethanediyl)] (9CI) (CA INDEX NAME)

RN 168788-11-2 HCA

CN Acetamide, 2,2,2-trifluoro-N-[3-[[4-[(trifluoroacetyl)amino]butyl]amino]propyl]-, mono(trifluoroacetate) (9CI) (CA INDEX NAME)

CM 1

CRN 158474-89-6 CMF C11 H17 F6 N3 O2

$$^{\rm O}_{||}$$
 $^{\rm O}_{||}$ $^{\rm O}_{||}$ $^{\rm O}_{||}$ $^{\rm O}_{||}$ $^{\rm F}_{3}$ $^{\rm C}$ $^{\rm C}$ $^{\rm NH}$ $^{\rm C}$ $^{\rm CH}_{2}$) $_{4}$ $^{\rm NH}$ $^{\rm C}$ $^{\rm CF}_{3}$

CM 2

CRN 76-05-1 CMF C2 H F3 O2

IC ICM C08G079-04

ICS A61K047-48; A61P021-06; A61P011-06; A61P009-10; A61P001-08; A61P035-00; A61P011-02; A61P001-12; A61P001-10

CC 63-6 (Pharmaceuticals)

Section cross-reference(s): 35

IT 83945-68-0P

(biodegradable polyphosphoramidates for controlled release of bioactive substances)

IT 100-36-7DP, N,N-Diethylethylenediamine, reaction products
with poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
108-00-9DP, N,N-Dimethylethylenediamine, reaction products
with poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
110-60-1DP, 1,4-Butanediamine, reaction products with
poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
15302-88-2DP, reaction products with poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
120527-66-4DP, reaction
products with polyamines 168788-11-2DP, reaction
products with poly(4-methyl-2-oxo-2-hydro-1,3,2-dioxaphospholane)
(biodegradable polyphosphoramidates for controlled release of bioactive substances)

L95 ANSWER 4 OF 19 HCA COPYRIGHT 2006 ACS on STN

Biodegradable polyphosphates for controlled release of drugs 137:389134 and genes and their preparation. Wang, Jun; Mao, Hai-Quan; Leong, Kam Weng (Johns Hopkins Singapore Pte. Ltd., Singapore). PCT Int. Appl. WO 2002092667 A1 20021121, 53 pp. DESIGNATED STATES: W: AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM; RW: AT, BE, BF, BJ, CF, CG, CH, CI, CM, CY, DE, DK, ES, FI, FR, GA, GB, GR, IE, IT, LU, MC, ML, MR, NE, NL, PT, SE, SN, TD, TG, TR. (English). CODEN: APPLICATION: WO 2002-SG90 20020514. PRIORITY: US PIXXD2. 2001-2001/PV290888 20010514.

AB The pos. chargeable polyphosphoester comprises .gtoreq.1 phosphoester linkage in the polymer backbone and .gtoreq.1 pos. chargeable group which is a substituent of a side chain attached to the polymer backbone through a phosphoester linkage. The

polyphosphoester is prepd. by polymg. .gtoreq.1 monomer to form a polymer with .gtoreq.1 phosphoester linkage in polymer backbone, reacting the polymer with a alc. with a chargeable group or its substituents. The compns. contg. the polyphosphoesters and biol. active substances are useful for delivery of drugs and genes. A controlled gene delivery system based on these polyphosphoesters is prepd. by complex coacervation of nucleic acid (DNA or RNA) with the polymers. The release rates can be manipulated by adjusting the charge ratios of polyphosphoesters to nucleic acids. This gene delivery system yields a higher gene expression in muscle when injected i.m.

IT 83906-57-4DP, chlorinated, esterified and hydrolyzed 83945-68-0DP, chlorinated, esterified and hydrolyzed

(prepn. of biodegradable polyphosphates for controlled release of drugs and genes)

RN 83906-57-4 HCA

CN Poly[oxyphosphinylideneoxy(1-methyl-1,2-ethanediyl)] (9CI) (CA INDEX NAME)

RN 83945-68-0 HCA

CN 1,3,2-Dioxaphospholane, 4-methyl-, 2-oxide, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 16352-26-4 CMF C3 H7 O3 P

TT 77987-49-6DP, Benzyl N-(2-hydroxyethyl)carbamate, reaction
products of chlorinated poly(4-methyl-2-hydro-1,3,2dioxaphospholane)

(prepn. of biodegradable polyphosphates for controlled release of drugs and genes)

RN 77987-49-6 HCA

CN Carbamic acid, (2-hydroxyethyl)-, phenylmethyl ester (9CI) (CA INDEX NAME)

O || Ph- CH₂- O- C- NH- CH₂- CH₂- OH

IC ICM C08G079-04

ICS A61K047-48; A61P021-06; A61P011-06; A61P009-10; A61P001-08; A61P035-00; A61P011-02; A61P001-12; A61P001-10

CC 63-5 (Pharmaceuticals)

83906-57-4DP, chlorinated, esterified and hydrolyzed 83945-68-0DP, chlorinated, esterified and hydrolyzed (prepn. of biodegradable polyphosphates for controlled release of drugs and genes)

TT 77987-49-6DP, Benzyl N-(2-hydroxyethyl)carbamate, reaction
products of chlorinated poly(4-methyl-2-hydro-1,3,2dioxaphospholane)

(prepn. of biodegradable polyphosphates for controlled release of drugs and genes)

L95 ANSWER 5 OF 19 HCA COPYRIGHT 2006 ACS on STN

- 135:348791 A Novel Biodegradable Gene Carrier Based on Polyphosphoester. Wang, Jun; Mao, Hai-Quan; Leong, Kam W. (Johns Hopkins, Singapore, 117597, Singapore). Journal of the American Chemical Society, 123(38), 9480-9481 (English) 2001. CODEN: JACSAT. ISSN: 0002-7863. Publisher: American Chemical Society.
- A new biodegradable gene carrier, poly(2-aminoethyl propylene AB phosphate) (PPE-EA) was designed and synthesized with relatively high mol. wt. A cytotoxicity study suggested the biocompatible nature of the polymer. Complexation between PPE-EA and plasmid protected DNA from nuclease degrdn. PPE-EA mediated a higher level of gene expression, but the transfection efficiency greatly depended on cell type. A unique feature of this system was the capability of controlled release of plasmid from PPE-EA/DNA complexes, achieved as a result of PPE-EA degrdn. The release-rate of plasmid could be adjusted by varying the charge ratio of PPE-EA to DNA. transfection activity of PPE-EA/DNA complexes was evaluated in HEK293 cells in comparison with poly(L-lysine) (PLL) and polyethylenimine (PEI). Transfection with PPE-EA/DNA complexes (charge ratios between 6 and 8) yielded 45-105-fold higher gene expression than PLL-mediated transfection. In the presence of chloroquine diphosphate (CQ, 100 M), a reagent known to disrupt endosomal membrane and enhance transfection of DNA complexes trafficked through the endo-lysosomal pathway, transfection efficiency was significantly increased.

IT 83906-57-4DP, aminoethylated

(prepn. and cytotoxicity of biodegradable gene carrier based on polyphosphoester)

RN 83906-57-4 HCA

CN Poly[oxyphosphinylideneoxy(1-methyl-1,2-ethanediyl)] (9CI) (CA INDEX NAME)

IT 77987-49-6, Benzyl N-(2-hydroxyethyl) carbamate

(prepn. and cytotoxicity of biodegradable gene carrier based on polyphosphoester)

RN 77987-49-6 HCA

CN Carbamic acid, (2-hydroxyethyl)-, phenylmethyl ester (9CI) (CA INDEX NAME)

IT 83906-57-4DP, (benzyloxycarbonyl) aminoethylated

(prepn. and cytotoxicity of biodegradable gene carrier based on polyphosphoester)

RN 83906-57-4 HCA

CN Poly[oxyphosphinylideneoxy(1-methyl-1,2-ethanediyl)] (9CI) (CA INDEX NAME)

CC 63-5 (Pharmaceuticals)

Section cross-reference(s): 3, 35

IT 83906-57-4DP, aminoethylated

(prepn. and cytotoxicity of biodegradable gene carrier based on polyphosphoester)

IT 16352-26-4 **77987-49-6**, Benzyl N-(2-hydroxyethyl) carbamate

(prepn. and cytotoxicity of biodegradable gene carrier based on polyphosphoester)

IT 83906-57-4DP, (benzyloxycarbonyl) aminoethylated

83906-57-4DP, chlorinated

(prepn. and cytotoxicity of biodegradable gene carrier based on polyphosphoester)

L95 ANSWER 6 OF 19 HCA COPYRIGHT 2006 ACS on STN

126:84180 Fixation of pharmacologically active amines on polyphosphonates. Part 3. Synthesis and preliminary in vitro cytotoxic studies of bis(2-chloroethyl)amino group containing polyphosphonates. Fontaine, Laurent; Marboeuf, Celine; Brosse, Jean Claude; Maingault, Martine; Dehaut, Frederic (Laboratoire Chimie Physicochimie Macromoleculaire, Universite Maine, Le Mans, F-72017, Fr.). Macromolecular Chemistry and Physics, 197(11), 3613-3621 (English) 1996. CODEN: MCHPES. ISSN: 1022-1352. Publisher: Huethiq & Wepf.

AB Water-sol. poly(3,6,9-trioxaundecamethylene phosphonate) with attached cytotoxic bis(2-chloroethyl)amine in the side chain was synthesized by chem. modification of the corresponding polyphosphonate via the Atherton-Todd reaction. The structure of the product was confirmed by 1H, 13C, and 31P NMR. Preliminary in vitro cytotoxic studies revealed that the polymeric deriv. exhibited an 8-fold cytotoxic potency against human hepatocellular carcinoma cell line HepG2 than against murine leukemia cell line L1210.

IT 124331-87-9DP, reaction product with bis(2-chloroethyl) amine 185537-34-2DP, reaction product with bis(2-chloroethyl)amine

(prepn. and cytotoxicity of bis(chloroethyl)amino-contg. polyphosphonate)

RN 124331-87-9 HCA

CN Poly(oxyphosphinylideneoxy-1,2-ethanediyloxy-1,2-ethanediyloxy-1,2-ethanediyloxy-1,2-ethanediyl) (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 185537-34-2 HCA

CN Phosphonic acid, dimethyl ester, polymer with 2,2'-[oxybis(2,1-ethanediyloxy)]bis[ethanol] (9CI) (CA INDEX NAME)

CM 1 868-85-9 CRN CMF C2 H7 O3 P MeO-PH-OMe CM 2 112-60-7 CRN CMF C8 H18 O5 CC 1-6 (Pharmacology) Section cross-reference(s): 35, 37, 38, 63 IT 124331-87-9DP, reaction product with bis(2-chloroethyl) amine 185537-34-2DP, reaction product with bis (2-chloroethyl) amine (prepn. and cytotoxicity of bis(chloroethyl)amino -contq. polyphosphonate) ANSWER 7 OF 19 HCA COPYRIGHT 2006 ACS on STN 125:277797 Adsorption of Bi3+ ion with diphenyl phosphonate-formaldehyde resin having tetraethylenepentamine. Mori, Hidehiko; Sekiguti, Masaki; Ochiai, Masanori; Watanabe, Makoto; Fujimura, Yoshikazu (Dep. of Industrial Chemistry, Chubu Univ., Kasugai, 487, Japan). Nippon Kagaku Kaishi (10), 917-919 (Japanese) 1996. CODEN: NKAKB8. ISSN: 0369-4577. Publisher: Nippon Kagakkai. AB Titled resin showed good adsorption ability for Bi3+ ion in hydrochloric acid soln., about 1.6 mmol per q resin based on a dry Bi3+ ion was adsorbed as anionic chloro complexes with the concn. > 0.1 mol dm-3, and the adsorption equil. was attained within Bi3+ chloro complex loaded on the resin could be readily eluted with 1.4 mol dm-3 nitric acid soln. IT 112-57-2D, Tetraethylenepentamine, reaction products with Di-Ph phosphonate-formaldehyde copolymer 68183-19-7D, Diphenyl phosphonate-formaldehyde copolymer, reaction products with tetraethylenepentamine

(adsorption of Bi3+ ion by di-Ph phosphonate-formaldehyde resin

with tetraethylenepentamine)

112-57-2 HCA

RN

1,2-Ethanediamine, N-(2-aminoethyl)-N'-[2-[(2-CN aminoethyl)amino]ethyl] - (9CI) (CA INDEX NAME) H₂N-CH₂-CH₂-NH-CH₂-CH₂-NH-CH₂-CH₂-NH-CH₂-CH₂-NH₂ RN 68183-19-7 HCA Phosphonic acid, diphenyl ester, polymer with formaldehyde (9CI) CN(CA INDEX NAME) CM 1 4712-55-4 CRN CMF C12 H11 O3 P O PhO-PH-OPh CM 2 CRN 50-00-0 CMF C H2 O $H_2C = 0$ CC 38-3 (Plastics Fabrication and Uses) Section cross-reference(s): 37 IT 112-57-2D, Tetraethylenepentamine, reaction products with Di-Ph phosphonate-formaldehyde copolymer 68183-19-7D, Diphenyl phosphonate-formaldehyde copolymer, reaction products with tetraethylenepentamine (adsorption of Bi3+ ion by di-Ph phosphonate-formaldehyde resin with tetraethylenepentamine) ANSWER 8 OF 19 HCA COPYRIGHT 2006 ACS on STN L95 123:144517 Amino acids coupled to poly(alkylene phosphates). Penczek, Stanislaw; Kaluzynski, Krzysztof; Baran, Jolanta (Cent. Mol. Macromol. Studies, Pol. Acad. Sci., Lodz, 90-363, Pol.). Macromol. 1992, Invited Lect. IUPAC Int. Symp. Macromol., 34th, Meeting Date 1992, 231-40. Editor(s): Kahovec, Jaroslav. VSP: Utrecht, Neth. (English) 1993. CODEN: 60KGAA. Methods are described of coupling amino acids or their esters to AB poly(alkylene phosphates). When, e.g. N,O-

bis(trimethylsilyl)glycine is used, its reaction with poly(alkylene

chlorophosphate) gives, after removing of the silyl blocking groups, poly(alkylene phosphates) with glycines attached to the phosphorus atoms via their nitrogens. The P-NH bonds hydrolyze in acid and close to neutral conditions faster than ester bonds. The rate consts. of hydrolysis depend on the pH of the medium and are 7.40 .times. 10-8s-1 and 2.40 .times. 10-8s-1 (pH 8.5-9.0) for the repeating unit amide and ester groups, resp. in poly(trimethylene phosphate) coupled to glycine. Poly(alkylene phosphates) are water-sol. nontoxic polymers considered as the drug carriers.

IT 59199-62-1D, chlorinated, reaction products with amino acid esters or dipeptides

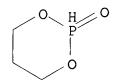
(prepn. and hydrolysis kinetics of amino acids coupled to poly(alkylene phosphates))

RN 59199-62-1 HCA

CN 1,3,2-Dioxaphosphorinane, 2-oxide, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 16352-21-9 CMF C3 H7 O3 P



IT 59199-86-9DP, chlorinated, reaction products with amino acid esters or dipeptides

(prepn. and hydrolysis kinetics of amino acids coupled to poly(alkylene phosphates))

RN 59199-86-9 HCA

CN Poly(oxyphosphinylideneoxy-1,3-propanediyl) (9CI) (CA INDEX NAME)

IT 159735-46-3DP, reaction products with amino acid esters

(prepn. and hydrolysis kinetics of **amino** acids coupled to poly(alkylene phosphates))

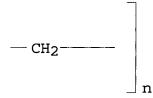
RN 159735-46-3 HCA

CN Poly[oxy(chlorophosphinylidene)oxy-1,2-ethanediyloxy-1,2-

ethanediyloxy-1,2-ethanediyloxy-1,2-ethanediyl] (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B



CC 34-2 (Amino Acids, Peptides, and Proteins)

Section cross-reference(s): 22, 63

ΙT 59199-62-1D, chlorinated, reaction products with amino acid esters or dipeptides

> (prepn. and hydrolysis kinetics of amino acids coupled to poly(alkylene phosphates))

59199-86-9DP, chlorinated, reaction products with ITamino acid esters or dipeptides

> (prepn. and hydrolysis kinetics of amino acids coupled to poly(alkylene phosphates))

627-74-7DP, reaction products with chlorinated poly(propylene IT phosphonate) 1738-68-7DP, reaction products with chlorinated poly(propylene phosphonate) 3082-75-5DP, Alanine ethyl ester, reaction products with chlorinated poly(propylene phosphonate) 6456-74-2DP, reaction products with chlorinated poly(propylene phosphonate) 16367-69-4DP, reaction products with chlorinated poly(propylene phosphonate) 17344-99-9DP, DL-Alanine ethyl ester, reaction products with chlorinated poly(propylene phosphonate) 30959-96-7DP, D-Alanine ethyl ester, reaction products with chlorinated poly(propylene phosphonate) 159735-46-3DP, reaction products with amino acid esters 166385-41-7DP, desilylated sodium salts

> (prepn. and hydrolysis kinetics of amino acids coupled to poly(alkylene phosphates))

L95 ANSWER 9 OF 19 HCA COPYRIGHT 2006 ACS on STN

122:10922 Amino acids attached to poly(alkylene phosphate)s. 1.
Synthesis. Kaluzynski, Krzysztof; Penczek, Stanislaw (Cent. Mol. Macromol. Studies, Acad. Sci., Lodz, 90-363, Pol.). Macromolecular Chemistry and Physics, 195(12), 3855-62 (English) 1994.
CODEN: MCHPES. ISSN: 1022-1352. Publisher: Huethig & Wepf.

AB Poly(alkylene phosphate)s with attached amino acids were synthesized by two methods for coupling of the amino acid moiety through P-NH bonds. Direct reaction of amino acid esters or of the N,O-bis(trimethylsilyl)-substituted amino acids with poly(alkylene chlorophosphate)s. The assumed microstructure of the resulting polymers was confirmed by anal. of the 1H, 13C and 31P NMR spectra. Preliminary data on the hydrolytic stability of amino acids modified poly(alkylene phosphate)s are also given.

IT 159735-46-3DP, reaction products with amino acid esters or dipeptides 159735-47-4DP, reaction products with amino acid esters or dipeptides

(prepn. and hydrolytic stability of degradable amino acid-coupled poly(alkylene phosphate)s)

RN 159735-46-3 HCA

CN Poly[oxy(chlorophosphinylidene)oxy-1,2-ethanediyloxy-1,2-ethanediyloxy-1,2-ethanediyloxy-1,2-ethanediyl) (9CI) (CA INDEX NAME)

PAGE 1-A

PAGE 1-B

RN 159735-47-4 HCA

CN Poly[oxy(chlorophosphinylidene)oxy-1,3-propanediyl] (9CI) (CA INDEX NAME)

CC 35-8 (Chemistry of Synthetic High Polymers)

IT 459-73-4DP, Ethyl glycinate, reaction products with poly(trialkylene phosphate chloride)s 627-74-7DP, reaction products with poly(trialkylene phosphate chloride)s 1738-68-7P, Benzyl glycinate 159735-46-3DP, reaction products with amino acid esters or dipeptides 159735-47-4DP, reaction products with amino acid esters or dipeptides

(prepn. and hydrolytic stability of degradable amino acid-coupled poly(alkylene phosphate)s)

L95 ANSWER 10 OF 19 HCA COPYRIGHT 2006 ACS on STN

118:42757 Rapid adsorption of gold(III) by phosphorus-containing chelate resins with polyamine. Mori, Hidehiko; Fujimura, Yoshikazu; Watanabe, Makoto; Sakurai, Makoto; Sinba, Hideharu (Coll. Eng., Chubu Univ., Kasugai, 487, Japan). Nippon Kagaku Kaishi (12), 1479-83 (Japanese) 1992. CODEN: NKAKB8. ISSN: 0369-4577.

The prepd. resins modified with triethylenetetramine, tetraethylenepentamine, and pentaethylenehexamine have the high adsorption capacity (2.3 mmol Au/g-resin). Gold(III) is readily adsorbed on the resins as anionic chloro-complexes from hydrochloric acid soln. Adsorption rate of gold(III) is high and the gold(III) was adsorbed effectively by passing gold(III) soln. (100 mg dm3) through the resin column at ion rate SV 600 h. The presence of 0.5 mol dm3 KH2PO4 at pH 4.2 or 0.2 mol dm3 HNO3 and the five-times amt. of Cu(II), and Al(III) to gold(III) at pH 0.5 do not affect the adsorption efficiency of gold(III). The gold(III) loaded on a column can be eluted with 0.5 mol dm3 HCl-50 vol.% acetone aq. soln. The Au-loaded resin which was treated with NaBH4 in ethanol can be utilized as a catalyst for hydrogenation of acetylene.

112-24-3D, Triethylenetetramine, reaction product with di-Ph phosphonate-formaldehyde copolymer 112-57-2D,
Tetraethylenepentamine, reaction product with di-Ph phosphonate-formaldehyde copolymer 4067-16-7D,
Pentaethylenehexamine, reaction product with di-Ph phosphonate-formaldehyde copolymer 68183-19-7D, Diphenyl phosphonate-formaldehyde copolymer, reaction products with polyamines

(adsorption by, of gold)

RN 112-24-3 HCA

CN 1,2-Ethanediamine, N,N'-bis(2-aminoethyl)- (9CI) (CA INDEX NAME)

 $H_2N-CH_2-CH_2-NH-CH_2-CH_2-NH-CH_2-CH_2-NH_2$ RN 112-57-2 HCA 1,2-Ethanediamine, N-(2-aminoethyl)-N'-[2-[(2-CN aminoethyl)amino]ethyl] - (9CI) (CA INDEX NAME) $H_2N-CH_2-CH_2-NH-CH_2-CH_2-NH-CH_2-CH_2-NH-CH_2-CH_2-NH_2$ RN4067-16-7 HCA CN 3,6,9,12-Tetraazatetradecane-1,14-diamine (9CI) (CA INDEX NAME) PAGE 1-A H₂N-CH₂-CH₂-NH-CH₂-CH₂-NH-CH₂-CH₂-NH-CH₂-CH₂-NH-CH₂-PAGE 1-B $-CH_2-NH_2$ RN 68183-19-7 HCA Phosphonic acid, diphenyl ester, polymer with formaldehyde (9CI) CN (CA INDEX NAME) CM 1 CRN 4712-55-4 CMF C12 H11 O3 P PhO-PH-OPh CM 2 CRN 50-00-0 CMF C H2 O

 $H_2C = 0$

- CC 54-2 (Extractive Metallurgy)
 Section cross-reference(s): 38, 45, 67
- IT 112-24-3D, Triethylenetetramine, reaction product with di-Ph
 phosphonate-formaldehyde copolymer 112-57-2D,
 Tetraethylenepentamine, reaction product with di-Ph
 phosphonate-formaldehyde copolymer 4067-16-7D,
 Pentaethylenehexamine, reaction product with di-Ph
 phosphonate-formaldehyde copolymer 68183-19-7D, Diphenyl
 phosphonate-formaldehyde copolymer, reaction products with
 polyamines

(adsorption by, of gold)

- L95 ANSWER 11 OF 19 HCA COPYRIGHT 2006 ACS on STN
- 112:81400 Adsorption and separation of gold(III) and platinum(IV) by phosphorus-containing chelate resin with tetraethylenepentamine groups. Mori, Hidehiko; Shinba, Hideharu; Fujimura, Yoshikazu; Takegami, Yoshinobu (Coll. Eng., Chubu Univ., Kasugai, 487, Japan). Nippon Kagaku Kaishi (11), 1855-8 (Japanese) 1989. CODEN: NKAKB8. ISSN: 0369-4577.
- AB A chelate resin was prepd. by the reaction of a di-Ph phosphonate-formaldehyde resin with tetraethylenepentamine for 4 h at 100-140.degree.. The adsorption capacity of the chelating resin of Au(III) and Pt(IV) was .apprx.2.3 mmol/g on a dry basis. rate of adsorption is high and the soln. passes effectively through the resin column. The chelating resin showed good adsorption ability for Au(III) and Pt(IV) at pH 1.6-6.0, and the selective sepn. of Au(III) and Pt(IV) from other di- and tri-valent metal ions is possible by HCl soln. The Pt(IV) and Au(III) ions loaded on a column are sepd. from each other by consecutive elution with 1.0 mol/L NaOH then with 2.5% KCN soln. The recovery of Au(III) and Pt(IV) from a large amt. of numerous metal ion samples, seawater, and wastewater was carried out and the rate of preconcn. for both ions was examd.
- 1T 112-57-2D, Tetraethylenepentamine, reaction product with
 di-Ph phosphonate-formaldehyde copolymer 68183-19-7D,
 Diphenyl phosphonate-formaldehyde copolymer, reaction product with
 tetraethylenepentamine

(extn. by, of gold and platinum)

RN 112-57-2 HCA

CN 1,2-Ethanediamine, N-(2-aminoethyl)-N'-[2-[(2-aminoethyl)amino]ethyl]- (9CI) (CA INDEX NAME)

 $H_2N-CH_2-CH_2-NH-CH_2-CH_2-NH-CH_2-CH_2-NH-CH_2-CH_2-NH_2$

RN 68183-19-7 HCA

CN Phosphonic acid, diphenyl ester, polymer with formaldehyde (9CI)

```
(CA INDEX NAME)
     CM
          1
         4712-55-4
     CRN
     CMF
          C12 H11 O3 P
     0
PhO-PH-OPh
     CM
          2
     CRN
         50-00-0
     CMF
         C H2 O
H_2C==0
```

CC 54-2 (Extractive Metallurgy)
IT 112-57-2D, Tetraethylenepentamine, reaction product with di-Ph phosphonate-formaldehyde copolymer 68183-19-7D,

Diphenyl phosphonate-formaldehyde copolymer, reaction product with tetraethylenepentamine

(extn. by, of gold and platinum)

L95 ANSWER 12 OF 19 HCA COPYRIGHT 2006 ACS on STN

109:231681 Preparation of a phosphorus-containing chelating resin and its selectivity for ionic adsorption. Mori, Hidehiko; Fujimura, Yoshikazu; Takegami, Yoshinobu; Ohashi, Tsugiya (Coll. Eng., Chubu Univ., Kasugai, 487, Japan). Nippon Kagaku Kaishi (9), 1633-5 (Japanese) 1988. CODEN: NKAKB8. ISSN: 0369-4577.

AB A chelating resin was prepd. by the reaction of di-Ph phosphonate-CH2O resin with triethylenetetramine for 4 h at .apprx.100.degree. The adsorption capacity of the resin for divalent metal ions was 1.8 Cu2+ mmol/g-R (dry resin) and for trivalent metal ions 0.80 Cr3+ mmol/g-R. Elution curves for Ca2+, Mg2+, Cu2+, Cd2+, Zn2+, Co2+, Ni2+ and Pb2+ were recorded by using 0.02 mol/L citric acid and 0.05 mol/L HCl. The sepn. of metal ions from each other and the recovery of Ca2+ and Mg2+ from large amts. of other divalent metal ions were facilitated. The resin also showed good adsorption of Cr(III) from buffer solns. of pH 5.0 .apprx. 7.0 and of Cr2O72- from buffer solns. of pH 2.0 .apprx. 7.0.

IT 112-24-3DP, Triethylenetetramine, reaction products with diphenylphosphonate-formaldehyde copolymer 68183-19-7DP, Diphenylphosphonate-formaldehyde copolymer, reaction products with

```
triethylenetetramine
        (prepn. and adsorption of metals by)
RN
     112-24-3 HCA
CN
     1,2-Ethanediamine, N,N'-bis(2-aminoethyl)- (9CI) (CA INDEX NAME)
H_2N-CH_2-CH_2-NH-CH_2-CH_2-NH-CH_2-CH_2-NH_2
RN
     68183-19-7 HCA
     Phosphonic acid, diphenyl ester, polymer with formaldehyde (9CI)
CN
     (CA INDEX NAME)
     CM
          1
     CRN
          4712-55-4
     CMF
          C12 H11 O3 P
     0
PhO-PH-OPh
     CM
     CRN
          50-00-0
     CMF
          C H2 O
H_2C==0
CC
     35-8 (Chemistry of Synthetic High Polymers)
IT
     112-24-3DP, Triethylenetetramine, reaction products with
     diphenylphosphonate-formaldehyde copolymer 68183-19-7DP,
     Diphenylphosphonate-formaldehyde copolymer, reaction products with
     triethylenetetramine
        (prepn. and adsorption of metals by)
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L95 ANSWER 13 OF 19 HCA COPYRIGHT 2006 ACS on STN

AB Poly(2-hydro-2-oxo-1,3,2-dioxaphosporinane) was quant. chlorinated, and the resulting polymer was reacted with excess of imidazole to

^{107:116051} Synthesis of poly(alkylene phosphate)s with N-containing bases in the side chains. II. 9-N-oxoethyleneadenine on the poly(trimethylene phosphate) chain. Lapienis, Grzegorz; Penczek, Stanislaw; Aleksyuk, G. P.; Kropachev, V. A. (Cent. Mol. Macromol. Stud., Pol. Acad. Sci., Lodz, 90-362, Pol.). Journal of Polymer Science, Part A: Polymer Chemistry, 25(7), 1729-36 (English) 1987. CODEN: JPACEC. ISSN: 0887-624X.

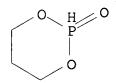
give a highly reactive polyesteramide. The polyesteramide was treated with N-hydroxyethyladenine with blocked amino group to give polyphosphates with N-oxyethyleneadenine in the side chains. The final polymers (no.-av. mol. wt. > 104) as well as the intermediate products were characterized by H-, C-13, and P-31 NMR spectroscopy. According to the spectra and anal. data quant. conversion was achieved.

RN 59199-62-1 HCA

CN 1,3,2-Dioxaphosphorinane, 2-oxide, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 16352-21-9 CMF C3 H7 O3 P



RN 59199-86-9 HCA

CN Poly(oxyphosphinylideneoxy-1,3-propanediyl) (9CI) (CA INDEX NAME)

CC 35-7 (Chemistry of Synthetic High Polymers) Section cross-reference(s): 36

TT 707-99-3DP, reaction products with chlorinated poly(oxodioxaphosphorinane)-imidazole reaction products 707-99-3P 59199-62-1DP, chlorinated, reaction products with imidazole and dimethylaminomethylene(hydroxyethyl)adenine or (hydroxyethyl)adenine 59199-86-9DP, chlorinated, reaction products with imidazole and dimethylaminomethylene (hydroxyethyl)adenine or (hydroxyethyl)adenine 110213-40-6DP, reaction products with chlorinated poly(oxodioxaphosphorinane)-

imidazole reaction products
 (prepn. and structure of)

L95 ANSWER 14 OF 19 HCA COPYRIGHT 2006 ACS on STN

97:128472 Studies on properties and reactivity of diphenyl phosphonate. XI. Preparation of a strongly basic anion exchanger based on diphenyl phosphonate-butyraldehyde resin. Mori, Hidehiko; Fujimura, Yoshikazu; Takegami, Yoshinobu (Dep. Ind. Chem., Chubu Inst. Technol., Kasugai, 487, Japan). Nippon Kagaku Kaishi (7), 1163-6 (Japanese) 1982. CODEN: NKAKB8. ISSN: 0369-4577.

AB A strongly basic anion exchange resin was prepd. by chloromethylation of an HCHO-crosslinked di-PH phosphonate-butyraldehyde resin, followed by amination of the ClCH2 group. The ion exchange capacity of the dry resin was 1.05 mequiv/g and the time to reach equil. was 10-15 min. This resin showed selective adsorption for CrO42-, MoO42-, and WO42- ions. These ions loaded on the resin were sepd. chromatog. from each other by sequential elution: Mo6+ was stripped by elution with a 1.6% NaCl soln. of pH 7.2, then Cr6+ with a 1.6% NaCl soln. of pH 7.8 and W6+ with 1.6% NaCl-0.1 mol/L NaOH, resp. The recoveries of CrO42-, MoO42- and WO42- ions at 103 mg/L each in mixed solns. of citric acid, sulfate, I-, and Cl- at 103 ppm were 100, 97, and 99%, resp.

IT 83105-25-3D, chloromethylated, aminated

(anion exchanger, for sepn. of molybdate and tungstate and chromate)

RN 83105-25-3 HCA

CN Phosphonic acid, diphenyl ester, polymer with butanal and formaldehyde (9CI) (CA INDEX NAME)

CM 1

CRN 4712-55-4 CMF C12 H11 O3 P

0 || PhO-- PH-- OPh

CM 2

CRN 123-72-8 CMF C4 H8 O

 $H_3C-CH_2-CH_2-CH=0$

CM 3

CRN 50-00-0 CMF C H2 O

 $H_2C = 0$

CC 37-3 (Plastics Manufacture and Processing)

IT 83105-25-3D, chloromethylated, aminated

(anion exchanger, for sepn. of molybdate and tungstate and chromate)

L95 ANSWER 15 OF 19 HCA COPYRIGHT 2006 ACS on STN
95:116597 Studies on property and reactivity of diphenyl phosphonate.
IX. Preparation of anion exchanger based on phosphorus-containing resin and its selective adsorption for anions. Mori, Hidehiko; Fujimura, Yoshikazu; Takegami, Yoshinobu (Dep. Ind. Chem., Chubu Inst. Technol., Kasugai, 487, Japan). Nippon Kagaku Kaishi (8), 1279-82 (Japanese) 1981. CODEN: NKAKB8. ISSN: 0369-4577.

AB The weakly basic anion exchange resin was prepd. by the chloromethylation of di-Ph phosphonate-formaldehyde condensation

chloromethylation of di-Ph phosphonate-formaldehyde condensation resin, followed by amination of the chloromethyl group. The ion exchange capacity of the dry resin was 0.50-1.35 mequiv/g, depending on the conditions of the amination. The optimum pH range for ion exchange was 2.0-6.0 and the time to reach an equil. state was 10-15 min. This resin showed selective adsorption for CrO42-, MoO42- and WO42- ions. Cr4+, Mo4+, and W4+ adsorbed on the resin could be completely eluted with 0.5 M NaOH soln. Using an acid soln. as an eluent, only Mo4+ could be stripped with 0.3 wt.% H2O2-1.5 M H2SO4 soln. The sepn. of Cr4+ and Mo4+ and the recovery of Cr4+ from the dil. soln. (10-3 ppm) were also investigated.

IT 68183-19-7D, chloromethylated, aminated (anion exchangers, selective adsorption of)

RN 68183-19-7 HCA

CN Phosphonic acid, diphenyl ester, polymer with formaldehyde (9CI) (CA INDEX NAME)

CM 1

CRN 4712-55-4 CMF C12 H11 O3 P

 $\begin{array}{c} \text{O} \\ \parallel \\ \text{PhO-PH-OPh} \end{array}$

CM 2

CRN 50-00-0 CMF C H2 O

 $H_2C = 0$

CC 37-3 (Plastics Fabrication and Uses)
IT 68183-19-7D, chloromethylated, aminated
(anion exchangers, selective adsorption of)

L95 ANSWER 16 OF 19 HCA COPYRIGHT 2006 ACS on STN
92:42899 Composition for cellular polyurethane. Sharafanov, V. T.;
Vinogradova, L. G.; Barynin, V. A.; Kaver, A. I.; Smyslov, V. I.;
Zubareva, L. S. (USSR). U.S.S.R. SU 696032 19791105
From: Otkrytiya, Izobret., Prom. Obraztsy, Tovarnye Znaki 1979,
(41), 100. (Russian). CODEN: URXXAF. APPLICATION: SU 1977-2522820
19770822.

AB Cellular polyurethanes with improved mech. strength and heat resistance were prepd. from a compn. consisting of 6.5-21.0 wt.% oxyethylated diphosphonate of diethylene glycol in addn. to polyisocyanate 49-62, oxypropylated ethylenediamine 19-29, surfactant 0.6-0.8, and blowing agent 1.8-11.3 wt.%.

RN 72441-53-3 HCA

CN Poly(oxy-1,2-ethanediyl), .alpha.,.alpha.'-(1,9-dioxido-2,5,8-trioxa-1,9-diphosphanonane-1,9-diyl)bis[.omega.-hydroxy-(9CI) (CA INDEX NAME)

PAGE 1-A

HO
$$CH_2 - CH_2 - O$$
 $PH - O - CH_2 - CH_2 - O - CH_2 - C$

PAGE 1-B

$$-$$
 PH $-$ O- CH₂- CH₂ OH

IC C08G018-14; C08L075-08

CC 36-6 (Plastics Manufacture and Processing)

IT 51178-86-0D, polymer with oxyethylated diethylene glycol diphosphonate and polyisocyanate **72441-53-3D**, polymer with oxypropylated **ethylenediamine** and polyisocyanate (cellular, heat-resistant)

L95 ANSWER 17 OF 19 HCA COPYRIGHT 2006 ACS on STN
78:30806 Organic phosphonate hardeners for epoxy resins. Oakes,
Vincent; Hutton, Ronald Eric; Cross, David F. W. (Interstab Ltd.)'.
Ger. Offen. DE 2212390 19720921, 22 pp. (German). CODEN:
GWXXBX. PRIORITY: GB 1971-7170 19710318.

The title hardeners, prepd. by condensation of polyamines with AB aldehydes and ketones, and then with di-O-substituted phosphite, give resins with lower flammability, better light stability, and better metal-resin adhesion. Thus an aquimolar amt. of diethylenetriamine is added portionwise at 30-40.deg, with stirring to paraformaldehyde, followed by an equimolar amt. of HOP(OMe)2. Heating to 80.deg./5 mm gives dimethyl diethylenetriaminomethanephos phonate (I) H2NCH2CH2NHCH2CH2NHCH2P(O)(OMe)2 [37114-84-4]. Bisphenol A diglycidyl ether (epoxy value 0.53/100 g resin) (100 g) mixed with 51 g I hardened at room temp., and then at 50.deg. for 48 hr to give a solid resin which is self-extinguishing in 6 sec. Other examples (47) are given, including hardeners prepd. from acetone, PhCHO, tetraethylenepentamine, diphenyl phosphite, ethylenediamine, p-phenylenediamine, and 4,4'diaminodiphenylmethane.

IT 39297-31-9 40623-48-1

(reaction with acetone and triethylenetetramine)

RN 39297-31-9 HCA

CN Poly[oxy(methyl-1,2-ethanediyl)], .alpha.-(ethoxyphosphinyl)-.omega.-[(ethoxyphosphinyl)oxy]- (9CI) (CA INDEX NAME)

RN 40623-48-1 HCA

CN Poly(oxy-1,2-ethanediyl), .alpha.-(ethoxyphosphinyl)-.omega.[(ethoxyphosphinyl)oxy]- (9CI) (CA INDEX NAME)

$$\begin{array}{c|c} \mathsf{C} & \mathsf{C} \\ || & \mathsf{C} \\ \mathsf{PH} & \mathsf{CH}_2 - \mathsf{CH}_2 - \mathsf{CH}_2 - \mathsf{CH}_2 \\ & \mathsf{D} \\ \end{array}$$

IT 107-15-3, reactions

(with diethyl phosphite and paraformaldehyde)

RN 107-15-3 HCA

CN 1,2-Ethanediamine (9CI) (CA INDEX NAME)

 $H_2N-CH_2-CH_2-NH_2$

IC C07F; C08G

CC 36-6 (Plastics Manufacture and Processing)

IT **39297-31-9** 40550-06-9 40550-07-0 40550-08-1 **40623-48-1**

(reaction with acetone and triethylenetetramine)

IT 107-15-3, reactions

(with diethyl phosphite and paraformaldehyde)

L95 ANSWER 18 OF 19 HCA COPYRIGHT 2006 ACS on STN
76:114359 Phosphorus-containing polymers. Santo, John E.; Vandegaer,
Jan E. (Pennwalt Corp.). U.S. US 3624248 19711130, 4 pp.
Division of U.S. 3,492,380 (CA 72;79882g). (English). CODEN:
USXXAM. APPLICATION: US 1970-403 19700106.

AB A P-contg. solid polymer which is sol. in alk. but not acid media is prepd. by interfacial polycondensation of p-(chlorocarbonyl)phenyl dichlorophosphate (I) [20070-01-3] and diamines or triamines and used to encapsulate liq., solid, or gas, e.g. insecticides. Thus, 3.1 g hexamethylenediamine in 200 ml water is agitated with a soln. of 5 g I in 5 ml benzene for a few min and the capsules are sepd. The p-(chlorocarbonyl)phenyl dichlorophosphate-hexamethylenediamine polymer [26680-16-0] encapsulates .sim.50 ml water. The polymer shells rupture in vacuum, are stable for >15 min at pH .leq.4 but burst in 2 min at pH 8.

IT 107-15-3, 1,2-Ethanediamine

(microcapsules from polymers contg., manuf. of, by interfacial polymn.)

RN 107-15-3 HCA

CN 1,2-Ethanediamine (9CI) (CA INDEX NAME)

 $H_2N-CH_2-CH_2-NH_2$

IT 26680-16-0P 26680-17-1P

(microcapsules of, manuf. of, by interfacial polymn.)

RN 26680-16-0 HCA

CN Phosphorodichloridic acid, 4-(chlorocarbonyl)phenyl ester, polymer with 1,6-hexanediamine (9CI) (CA INDEX NAME)

CM 1

CRN 20070-01-3 CMF C7 H4 Cl3 O3 P

CM 2

CRN 124-09-4 CMF C6 H16 N2

 $H_2N-(CH_2)_6-NH_2$

RN 26680-17-1 HCA

CN Phosphorodichloridic acid, 4-(chlorocarbonyl)phenyl ester, polymer with piperazine (9CI) (CA INDEX NAME)

CM 1

CRN 20070-01-3 CMF C7 H4 Cl3 O3 P

CM 2

CRN 110-85-0 CMF C4 H10 N2

IC C08G

INCL 260047000CZ

37 (Plastics Fabrication and Uses)

Section cross-reference(s): 25

107-15-3, 1,2-Ethanediamine 9016-87-9, IT 111-40-0 Isocyanic acid, polymethylenepolyphenylene ester 24802-03-7 (microcapsules from polymers contg., manuf. of, by interfacial polymn.)

IT26680-16-0P 26680-17-1P

(microcapsules of, manuf. of, by interfacial polymn.)

ANSWER 19 OF 19 HCA COPYRIGHT 2006 ACS on STN Encapsulation with polymer material. Santo, John E.; Vandegaer, Jan E. (Pennwalt Corp.). U.S. US 3492380 **19700127**, 3 pp. (English). CODEN: USXXAM. APPLICATION: US 1967-621135 19670307.

AB Solid or liq. particles are encapsulated by an interfacial polycondensation reaction with a P-contq. polymer shell, which is acid resistant, but dissolves in bases. Thus, 5 g p-(chlorocarbonyl)phenyl reaction with a P-contg. polymer shell, which is acid resistant, but dissolves in bases. Thus, 5 q p-(chlorocarbonyl)phenyl dichlorophosphate (I) (prepd. by heating p-hydroxybenzoic acid with PCl5) in 5 ml C6H6 was added to an agitated soln. of 3.1 g hexamethylenediamine in 200 ml H2O to form solid shells, which ruptured under vacuum and released .apprx.50ml

H2O. The shells were stable for 15 min in aq. solns. of pH <7, burst after 12 min at pH 7, and burst after 2 min at pH>7. A mixt. of I, polymethylene poly(phenylisocyanate) (PAPI), sebacoyl chloride, poly(vinyl alc.) and Thimet was added to a soln. of ethylenediamine, diethylenetriamine, and Na2CO3 to form an encapsulated insecticide. 2,2-Bis(4-hydroxyphenyl)propane bis-(dichlorophosphate), prepd. by adding PCl3 to a mixt. of bisphenol A and POCl3, was similarly used to encapsulate Diazinon. The use of p-(chlorocarbonylmethylene)phenyl dichlorophosphate instead of I is claimed.

IT 26680-16-0 26680-17-1

(Diazinon encapsulation with)

RN 26680-16-0 HCA

CN Phosphorodichloridic acid, 4-(chlorocarbonyl)phenyl ester, polymer with 1,6-hexanediamine (9CI) (CA INDEX NAME)

CM 1

CRN 20070-01-3 CMF C7 H4 Cl3 O3 P

CM 2

CRN 124-09-4 CMF C6 H16 N2

 H_2N^- (CH₂)₆-NH₂

RN 26680-17-1 HCA

CN Phosphorodichloridic acid, 4-(chlorocarbonyl)phenyl ester, polymer with piperazine (9CI) (CA INDEX NAME)

CM 1

CRN 20070-01-3 CMF C7 H4 Cl3 O3 P

CRN 110-85-0 CMF C4 H10 N2

IT 107-15-3, Ethylenediamine

(polymers with (chloroformyl) phenyl phosphorodichloridate and isocyanic acid esters, Thimet encapsulation with)

RN 107-15-3 HCA

CN 1,2-Ethanediamine (9CI) (CA INDEX NAME)

 $H_2N-CH_2-CH_2-NH_2$

IC A61J; B29C

INCL 264004000

CC 36 (Plastics Manufacture and Processing)

IT 111-19-3D, Sebacoyl chloride, polymers with diethylenetriamine and isocyanic acid esters 26680-16-0 26680-17-1

(Diazinon encapsulation with)

IT 107-15-3, Ethylenediamine 111-40-0

(polymers with (chloroformyl) phenyl phosphorodichloridate and isocyanic acid esters, Thimet encapsulation with)

=> d 196 1-47 ti

L96 ANSWER 1 OF 47 HCA COPYRIGHT 2006 ACS on STN

TI Poly(arylazophosphonate)s: new arylazophosphonate-containing monomers for synthesis of laser-structurable polymers

- L96 ANSWER 2 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Imide-amide rearrangement of cyclic phosphorimidates: a mechanistic study
- L96 ANSWER 3 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Synthesis of new phosphonic derivatives bearing fluorinated chains
- L96 ANSWER 4 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Arylazophosphonate containing polymers designed for XeCl excimer laser ablation
- L96 ANSWER 5 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Laser ablation of arylazo-containing polymers
- L96 ANSWER 6 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Preparation of oligo-amides and polyamide which contain the 4-dialkylamino-4'-(alkylsulfonyl)azobenzene chromophore
- L96 ANSWER 7 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Photochemical properties of poly(arylazophosphonates)
- L96 ANSWER 8 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Poly(arylazophosphonate)s: A New Polymer Class with Arylazophosphonate Units in the Main Chain
- L96 ANSWER 9 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Synthesis and MRI enhancement of iron(III) complexes with phosphate-EDTA-polyglycol terpolymer ligands
- L96 ANSWER 10 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Multilayer recording element suitable for production of flexographic printing plate by digital information transmission
- L96 ANSWER 11 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Manufacture of linear polyphosphazenes
- L96 ANSWER 12 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Polymers with arylazophosphonate units. Application for laser ablation
- L96 ANSWER 13 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Silver halide photographic material with storage stability
- L96 ANSWER 14 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Preparation of novel phosphoramidate and phophorothioamidate oligomeric compounds

- L96 ANSWER 15 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Coating compositions containing phosphinylphosphoimides
- L96 ANSWER 16 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI The synthesis of phosphinylphosphorimidic hydroxyethyl acrylate and the electrical properties of its polymer produced by ultraviolet-irradiation-induced polymerization
- L96 ANSWER 17 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Preparation of phosphazene oligomers
- L96 ANSWER 18 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Wood-impregnating resin compositions with improved fire and weather resistance
- L96 ANSWER 19 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Synthesis and antitumor activity of polyphosphates containing both nitrogen mustard and lipophilic groups
- L96 ANSWER 20 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Phosphorus-containing thiuram disulfides as chain-transfer agents in the preparation of fire-resistant polymers
- L96 ANSWER 21 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Phosphorus functional telomers of vinyl polymers through functional thermal iniferter
- L96 ANSWER 22 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Thermal functional polymeric "iniferters" based on phosphorus-containing poly(thiuram disulfides); synthesis and characterization
- L96 ANSWER 23 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Syntheses of phosphorus based flame retardant polymers
- L96 ANSWER 24 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Styrene copolymers
- L96 ANSWER 25 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Biocide polymers. III. Synthesis of polyurethanes, polyureas, and their thioanalogs having phosphoric acid-phenyl ester diamide structural units
- L96 ANSWER 26 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Fire-resistant urethane-modified isocyanurate foams
- L96 ANSWER 27 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Crosslinked, carboxylic polymers suitable as thickeners

- L96 ANSWER 28 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Fire-resistant polyester fibers
- L96 ANSWER 29 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Heat resistant polymers of oxidized styrylphosphine
- L96 ANSWER 30 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Heat resistant polymers of oxidized styrylphosphine
- L96 ANSWER 31 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Heat resistant polymers of oxidized styrylphosphine
- L96 ANSWER 32 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Textiles fire-retardant treated with copolycondensed vinylphosphonates
- L96 ANSWER 33 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Aniline-based reaction product
- L96 ANSWER 34 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Flame-resistant polyamides
- L96 ANSWER 35 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI On polymerization of organofluorine cyclic amidophosphates
- L96 ANSWER 36 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Polyamides from fluoroalkyl dichlorophosphates
- L96 ANSWER 37 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Fire-resistant polyester fibers
- L96 ANSWER 38 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Phosphor-containing polysulfones
- L96 ANSWER 39 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Electric birefringence, conformation, and rigidity of the molecules of cellulose esters in solutions
- L96 ANSWER 40 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Pharmacologically active polymers. 7. Cyclophosphamide and steroid hormone containing polymers as potential anticancer compounds
- L96 ANSWER 41 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Copolycondensed vinyl phosphonates as fire retardants
- L96 ANSWER 42 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Bis (azidophosphoranes)

- L96 ANSWER 43 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Polymers containing boron, oxygen and phosphorus
- L96 ANSWER 44 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Polymers prepared from cyclic phosphite esters and p-benzoquinones
- L96 ANSWER 45 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Poly(tripropylene glycol phosphoramidates), useful as flameproofing agents
- L96 ANSWER 46 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI High-molecular-weight crosslinked polyurethane resins
- L96 ANSWER 47 OF 47 HCA COPYRIGHT 2006 ACS on STN
- TI Adducts of polyalkene wax and alkyl phosphite

=> d 196 9,18,25,39,40 cbib abs hitstr hitind

- L96 ANSWER 9 OF 47 HCA COPYRIGHT 2006 ACS on STN
- 129:54667 Synthesis and MRI enhancement of iron(III) complexes with phosphate-EDTA-polyglycol terpolymer ligands. Yu, Kaichao; Feng, Jiwen; Hu, Hongbing; Lu, Guang; Zhuo, Renxi; Li, Liyun; Ye, Chaohui (Wuhan Inst. Physics Mathematics, Chinese Acad. Scis., Wuhan, 430071, Peop. Rep. China). Bopuxue Zazhi, 15(2), 123-129 (Chinese) 1998. CODEN: BOZAE2. ISSN: 1000-4556. Publisher: Zhongguo Kexueyuan Wuhan Wuli Yanjiuso.
- AB A new series of terpolymer ligands were synthesized by the copolymn. of ethylene diamine tetraacetic dianhydride (EDTAA), Et phosphodichloridate (or Me phosphodichloridate, Bu phosphodichloridate) with diethylene glycol or triethylene glycol, and their Fe(III) complexes were prepd. The structures of these new ligands and Fe(III) complexes were confirmed by 1H NMR, IR and elemental anal. The longitudinal relaxation measurements on these macromol. complexes indicated that most of them have higher relativity as compared to Fe(III)-EDTA. The acute toxicity test using one of these Fe(III) complexes was performed and no obvious toxicity was obsd. T1-weighted contrast imaging of rat liver showed clear MRI enhancement after injection the new complex.
- 208517-82-2P, Diethylene glycol-ethylenediamine tetraacetic dianhydride-methyl phosphodichloridate copolymer (synthesis, structure, magnetic relaxation, toxicity, and MRI enhancement of iron(III) complexes with phosphate-EDTA-polyglycol terpolymer ligands)
- RN 208517-82-2 HCA
- CN Phosphorodichloridic acid, methyl ester, polymer with 4,4'-(1,2-ethanediyl)bis[2,6-morpholinedione] and

2,2'-oxybis[ethanol] (9CI) (CA INDEX NAME)

CM 1

CRN 23911-25-3 CMF C10 H12 N2 O6

CM 2

CRN 677-24-7 CMF C H3 Cl2 O2 P

CM 3

CRN 111-46-6 CMF C4 H10 O3

 ${\tt HO-CH_2-CH_2-O-CH_2-CH_2-OH}$

CC 35-5 (Chemistry of Synthetic High Polymers)
 Section cross-reference(s): 63
IT 208517-82-2P, Diethylene glycol-ethylenediamine

tetraacetic dianhydride-methyl phosphodichloridate copolymer 208517-83-3P 208517-84-4P 208517-85-5P 208517-86-6P 208517-87-7P

(synthesis, structure, magnetic relaxation, toxicity, and MRI enhancement of iron(III) complexes with phosphate-EDTA-polyglycol terpolymer ligands)

L96 ANSWER 18 OF 47 HCA COPYRIGHT 2006 ACS on STN

- 115:184630 Wood-impregnating resin compositions with improved fire and weather resistance. Nakahara, Makoto; Yoshiga, Ikuo; Izumi, Masao (Sanken Kako Co., Ltd., Japan). Jpn. Kokai Tokkyo Koho JP 03097502 A2 19910423 Heisei, 3 pp. (Japanese). CODEN: JKXXAF. APPLICATION: JP 1989-236094 19890912.
- Title compns. comprise unsatd. polyesters obtained from halo- or P-contg. aliph. polyols and unsatd. dicarboxylic acids or their anhydrides, and optionally aliph. dicarboxylic acids, acrylic compds., P-contg. plasticizers, and curing catalysts. Thus, a white ash specimen impregnated with a mixt. of adipic acid-dibromoneopentyl glycol-maleic anhydride copolymer, tris(2,3-dibromopropyl) phosphate, Me methacrylate, and benzoyl peroxide and cured at 140.degree. showed good wood pattern, Barcol hardness (GYZJ 935) 70, O index .gtoreq.40, and good weather resistance.

IT 136654-26-7P

(prepn. of, contg. acrylate esters and phosphate plasticizers and curing catalysts, impregnants for wood, fire- and weather-resistant)

RN 136654-26-7 HCA

CN Hexanedioic acid, polymer with bis(1-methylethyl) bis(2-hydroxyethyl)phosphoramidate, (2E)-2-butenedioic acid and methyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 54480-56-7 CMF C10 H24 N O5 P

$$\begin{array}{c} \text{i-PrO-} \overset{\text{O}}{\mid\mid} \\ \text{i-PrO-} \overset{\text{P-}}{\text{OPr-i}} \\ \mid \\ \text{HO-} \text{CH}_2\text{-} \text{CH}_2\text{-} \text{N-} \text{CH}_2\text{-} \text{CH}_2\text{-} \text{OH} \end{array}$$

CM 2

CRN 124-04-9

CMF C6 H10 O4

 $HO_2C-(CH_2)_4-CO_2H$

CM 3

CRN 110-17-8 CMF C4 H4 O4

Double bond geometry as shown.

CM 4

CRN 80-62-6 CMF C5 H8 O2

IC ICM B27K003-50

CC 37-6 (Plastics Manufacture and Processing) Section cross-reference(s): 35, 43

IT 136654-26-7P

(prepn. of, contg. acrylate esters and phosphate plasticizers and curing catalysts, impregnants for wood, fire- and weather-resistant)

L96 ANSWER 25 OF 47 HCA COPYRIGHT 2006 ACS on STN 94:157338 Biocide polymers. III. Synthesis of polyurethanes, polyureas, and their thioanalogs having phosphoric acid-phenyl ester diamide structural units. Hartmann, Manfred; Geschwend, G.; Klemm, D. (Sekt. Chem., Friedrich-Schiller-Univ. Jena, Jena, DDR-6900, Ger. Dem. Rep.). Acta Polymerica, 32(1), 24-6 (German) 1981. CODEN: ACPODY. ISSN: 0323-7648.

AB The soln. polymn. of PhOPO(NCS)2 [21049-95-6] (prepd. from PhOPOCl2 [770-12-7] and NH4SCN in C6H6-MeCN) with glycols and with alkanediamines gave polythiourethanes and polythioureas, resp. The mol. wts. of the latter were distinctly higher. No transitions were shown by the polymers at .ltoreq.200.degree. in differential

scanning calorimetry. Similar polymns. of PhOPO(NCO)2 [prepd. by condensing PhOPCl2 with KOCN and oxidizing PhOP(NCO)2 with SO2Cl2] gave polyurethanes and polyureas. The mol. wts. of the polyureas were comparable to those of the polythioureas.

IT 77223-94-0P 77223-95-1P 77223-96-2P

77223-97-3P 77223-98-4P 77223-99-5P

77224-00-1P 77224-01-2P 77224-02-3P

77224-03-4P 77224-04-5P 77224-05-6P

77224-06-7P

(prepn. of)

RN 77223-94-0 HCA

CN Phosphorodiisocyanatidic acid, phenyl ester, polymer with 1,2-ethanediol (9CI) (CA INDEX NAME)

CM 1

CRN 1844-12-8 CMF C8 H5 N2 O4 P

CM 2

CRN 107-21-1 CMF C2 H6 O2

 $HO-CH_2-CH_2-OH$

RN 77223-95-1 HCA

CN Phosphorodiisocyanatidic acid, phenyl ester, polymer with 1,4-butanediol (9CI) (CA INDEX NAME)

CM 1

CRN 1844-12-8 CMF C8 H5 N2 O4 P

CRN 110-63-4 CMF C4 H10 O2

 ${\tt HO^-}$ (CH₂)₄ $-{\tt OH}$

RN 77223-96-2 HCA

CN Phosphorodiisocyanatidic acid, phenyl ester, polymer with 2,2'-oxybis[ethanol] (9CI) (CA INDEX NAME)

CM 1

CRN 1844-12-8 CMF C8 H5 N2 O4 P

CM 2

CRN 111-46-6 CMF C4 H10 O3

 ${\tt HO-CH_2-CH_2-O-CH_2-CH_2-OH}$

RN 77223-97-3 HCA

CN Phosphorodiisocyanatidic acid, phenyl ester, polymer with 2,2'-[1,2-ethanediylbis(oxy)]bis[ethanol] (9CI) (CA INDEX NAME)

CM 1

CRN 1844-12-8 CMF C8 H5 N2 O4 P

CM 2

CRN 112-27-6 CMF C6 H14 O4

 ${\tt HO-CH_2-CH_2-O-CH_2-CH_2-O-CH_2-OH}$

RN 77223-98-4 HCA

CN Phosphorodiisocyanatidic acid, phenyl ester, polymer with 1,2-ethanediamine (9CI) (CA INDEX NAME)

CM 1

CRN 1844-12-8 CMF C8 H5 N2 O4 P

CM 2

CRN 107-15-3 CMF C2 H8 N2

 $H_2N-CH_2-CH_2-NH_2$

RN 77223-99-5 HCA

CN Phosphorodiisocyanatidic acid, phenyl ester, polymer with 1,6-hexanediamine (9CI) (CA INDEX NAME)

CRN 1844-12-8 CMF C8 H5 N2 O4 P

CM 2

CRN 124-09-4 CMF C6 H16 N2

 $H_2N-(CH_2)_6-NH_2$

RN 77224-00-1 HCA

CN Phosphorodi(isothiocyanatidic) acid, phenyl ester, polymer with 1,2-ethanediol (9CI) (CA INDEX NAME)

CM 1

CRN 21049-95-6 CMF C8 H5 N2 O2 P S2

CM 2

CRN 107-21-1 CMF C2 H6 O2

 $HO-CH_2-CH_2-OH$

RN 77224-01-2 HCA

CN Phosphorodi(isothiocyanatidic) acid, phenyl ester, polymer with

1,4-butanediol (9CI) (CA INDEX NAME)

CM 1

CRN 21049-95-6

CMF C8 H5 N2 O2 P S2

CM 2

CRN 110-63-4

CMF C4 H10 O2

$$^{\rm HO-}$$
 (CH₂)₄ $^{-}$ OH

RN 77224-02-3 HCA

CN Phosphorodi(isothiocyanatidic) acid, phenyl ester, polymer with 2,2'-oxybis[ethanol] (9CI) (CA INDEX NAME)

CM 1

CRN 21049-95-6

CMF C8 H5 N2 O2 P S2

CM 2

CRN 111-46-6

CMF C4 H10 O3

 $HO-CH_2-CH_2-O-CH_2-CH_2-OH$

RN 77224-03-4 HCA

CN Phosphorodi(isothiocyanatidic) acid, phenyl ester, polymer with 2,2'-[1,2-ethanediylbis(oxy)]bis[ethanol] (9CI) (CA INDEX NAME)

CM 1

CRN 21049-95-6 CMF C8 H5 N2 O2 P S2

CM 2

CRN 112-27-6 CMF C6 H14 O4

$${\tt HO-CH_2-CH_2-O-CH_2-CH_2-O-CH_2-CH_2-OH}$$

RN 77224-04-5 HCA

CN Phosphorodi(isothiocyanatidic) acid, phenyl ester, polymer with 1,2-ethanediamine (9CI) (CA INDEX NAME)

CM 1

CRN 21049-95-6 CMF C8 H5 N2 O2 P S2

$$S = C = N - P - N = C = S$$

CM 2

CRN 107-15-3 CMF C2 H8 N2 $H_2N-CH_2-CH_2-NH_2$

RN 77224-05-6 HCA

CN Phosphorodi(isothiocyanatidic) acid, phenyl ester, polymer with 1,4-butanediamine (9CI) (CA INDEX NAME)

CM 1

CRN 21049-95-6 CMF C8 H5 N2 O2 P S2

$$\begin{array}{c} OPh \\ | \\ P-N \longrightarrow C \longrightarrow S \\ | \\ O \end{array}$$

CM 2

CRN 110-60-1 CMF C4 H12 N2

 $H_2N-(CH_2)_4-NH_2$

RN 77224-06-7 HCA

CN Phosphorodi(isothiocyanatidic) acid, phenyl ester, polymer with 1,6-hexanediamine (9CI) (CA INDEX NAME)

CM 1

CRN 21049-95-6 CMF C8 H5 N2 O2 P S2

CM 2

CRN 124-09-4 CMF C6 H16 N2

 $H_2N-(CH_2)_6-NH_2$

CC 35-3 (Synthetic High Polymers) IT77210-99-2P 21049-95-6P 77211-00-8P 77211-01-9P 77211-02-0P 77211-06-4P 77211-03-1P 77211-04-2P 77211-05-3P 77211-07-5P 77211-08-6P 77211-09-7P 77211-10-0P 77211-11-1P 77223-94-0P 77223-95-1P 77223-96-2P 77223-97-3P 77223-98-4P 77223-99-5P 77224-00-1P 77224-01-2P 77224-02-3P 77224-03-4P 77224-04-5P 77224-05-6P 77224-06-7P (prepn. of)

L96 ANSWER 39 OF 47 HCA COPYRIGHT 2006 ACS on STN
82:86758 Electric birefringence, conformation, and rigidity of the
 molecules of cellulose esters in solutions. Lavrenko, P. N.;
 Ryumtsev, E. I.; Shtennikova, I. N.; Andreeva, L. N.; Pogodina, N.
 V.; Tsvetkov, V. N. (Inst. High Mol. Compounds, Leningrad, USSR).
 Journal of Polymer Science, Polymer Symposia, 44 (Organ. Struct.
 Polym. Solutions Gels, Microsymp. Macromol., 12th, 1973), 217-35
 (English) 1974. CODEN: JPYCAQ. ISSN: 0360-8905.

AB Solns. of several cellulose [9004-34-6] esters in dioxane showed high equil.and kinetic chain rigidity. The coincidence in signs of the Kerr and Maxwell effects in solns. of the samples indicated gen. relations shown by all rigid-chain polymers for which the longitudinal geometric axis of the mol. is also the symmetry axis of its polar and optical properties. For example, increasing the mol. wt. of cellulose carbanilate (I) [37251-21-1] in soln. changed the conformation of I mols. from a faintly-curved rod to a rigid Gaussian coil. In the Gaussian region (high mol. wt.) the Kerr effect depended on the longitudinal (with respect to the chain) component of the dipole moment formed by the C-O bonds forming a portion of the glucoside ring.

IT 54579-30-5

(elec. and flow birefringence of, in dioxane, detn. of)

RN 54579-30-5 HCA

CN Cellulose, (diphenoxyphosphinyl)phenylcarbamate (9CI) (CA INDEX NAME)

CM 1

CRN 168611-97-0 CMF C19 H16 N O5 P

CRN 9004-34-6 CMF Unspecified

CCI PMS, MAN

*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

CC 35-5 (Synthetic High Polymers) Section cross-reference(s): 69

IT 9004-34-6D, Cellulose, deriv., esters 9004-36-8 9004-57-3 9015-11-6 9032-47-7 9069-20-9 37251-21-1 **54579-30-5** (elec. and flow birefringence of, in dioxane, detn. of)

L96 ANSWER 40 OF 47 HCA COPYRIGHT 2006 ACS on STN 82:43784 Pharmacologically active polymers. 7. Cyclophosphamide and steroid hormone containing polymers as potential anticancer compounds. Batz, Hans G.; Ringsdorf, Helmut; Ritter, Helmut (Inst. Org. Chem., Univ. Mainz, Mainz, Fed. Rep. Ger.). Makromolekulare Chemie, 175(8), 2229-39 (English) 1974. CODEN: MACEAK. ISSN: 0025-116X.

AB Several acrylic and methacrylic esters and N-vinyl monomers of cyclophosphamide and testosterone derivs. were prepd. and homopolymd. or copolymd. with hydrophilic monomers to give water-sol. polymers. For example, 2-[N-(2-chloroethyl)-N-(3-.beta.-chloroethyl-2-oxoperhydro-1,3,2.lambda.5-oxazaphosphorin-2-yl)amino]ethyl methacrylate-2-methacryloyloxyethyltrimethylammonium chloride polymer [53899-92-6] was prepd. and showed pharmacol. activity against Yoshida ascites sarcoma in mice (5 mg/kg body wt.).

IT 53899-92-6

(pharmacol. activity of)

RN 53899-92-6 HCA

CN Ethanaminium, N,N,N-trimethyl-2-[(2-methyl-1-oxo-2-propenyl)oxy]-, chloride, polymer with 2-[(2-chloroethyl)[3-(2-chloroethyl)tetrahydro-2-oxido-2H-1,3,2-oxazaphosphorin-2-yl]amino]ethyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 53804-58-3 CMF C13 H23 Cl2 N2 O4 P

CRN 5039-78-1 CMF C9 H18 N O2 . Cl

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ || & || & || \\ \text{Me}_3\text{+N}-\text{CH}_2-\text{CH}_2-\text{O}-\text{C}-\text{C}-\text{Me} \end{array}$$

● cl - .

IT 53804-59-4P 53804-61-8P 53804-63-0P 53804-65-2P 53804-66-3P 53804-67-4P 53804-68-5P 53899-91-5P (prepn. of)

53804-59-4 HCA

CN 2-Propenoic acid, 2-methyl-, 2-[(2-chloroethyl)[3-(2-chloroethyl)tetrahydro-2-oxido-2H-1,3,2-oxazaphosphorin-2-yl]amino]ethyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

RN

CRN 53804-58-3 CMF C13 H23 Cl2 N2 O4 P

RN 53804-61-8 HCA

CN 2-Propenoic acid, 2-methyl-, 3-[(2-chloroethyl)[3-(2-chloroethyl)tetrahydro-2-oxido-2H-1,3,2-oxazaphosphorin-2-yl]amino]propyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 53804-60-7

CMF C14 H25 Cl2 N2 O4 P

RN 53804-63-0 HCA

CN 2-Propenoic acid, 2-methyl-, 3-[2-[bis(2-chloroethyl)amino]dihydro-2-oxido-2H-1,3,2-oxazaphosphorin-3(4H)-yl]propyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 53804-62-9

CMF C14 H25 Cl2 N2 O4 P

$$\begin{array}{c|c} & \text{O} & \text{CH}_2 \\ & \parallel & \parallel \\ & \text{(CH}_2)_3 - \text{O} - \text{C} - \text{C} - \text{Me} \\ & \mid & \text{O} \\ & \parallel & \parallel \\ & \text{O} \\ & N - \text{CH}_2 - \text{CH}_2 \text{Cl} \\ & \mid & \parallel \\ & \text{O} \\ & \text{CH}_2 - \text{CH}_2 \text{Cl} \\ \end{array}$$

RN 53804-65-2 HCA

CN 2-Propenoic acid, 3-[2-[bis(2-chloroethyl)amino]dihydro-2-oxido-2H-1,3,2-oxazaphosphorin-3(4H)-yl]propyl ester, homopolymer (9CI) (CA INDEX NAME)

CM 1

CRN 53804-64-1 CMF C13 H23 Cl2 N2 O4 P

RN 53804-66-3 HCA

CN 2-Propenoic acid, 2-methyl-, 2-[(2-chloroethyl)[3-(2-chloroethyl)tetrahydro-2-oxido-2H-1,3,2-oxazaphosphorin-2-yl]amino]ethyl ester, polymer with 2-(methylsulfinyl)ethyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 53804-58-3 CMF C13 H23 Cl2 N2 O4 P

CM 2

CRN 14794-09-3 CMF C7 H12 O3 S

RN 53804-67-4 HCA

CN 2-Propenoic acid, 3-[2-[bis(2-chloroethyl)amino]dihydro-2-oxido-2H-1,3,2-oxazaphosphorin-3(4H)-yl]propyl ester, polymer with

2-(methylsulfinyl)ethyl 2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 53804-64-1

CMF C13 H23 Cl2 N2 O4 P

$$(CH_{2})_{3}-O-C-CH=CH_{2}$$

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CM 2

CRN 42356-51-4 CMF C6 H10 O3 S

$$\begin{array}{c} \text{O} & \text{O} \\ || & || \\ \text{Me-S-CH}_2\text{-CH}_2\text{-O-C-CH----} \text{CH}_2 \end{array}$$

RN 53804-68-5 HCA

CN Carbamic acid, ethenyl-, 3-[2-[bis(2-chloroethyl)amino]dihydro-2-oxido-2H-1,3,2-oxazaphosphorin-3(4H)-yl]propyl ester, polymer with 1-ethenyl-2-pyrrolidinone (9CI) (CA INDEX NAME)

CM 1

CRN 53803-76-2

CMF C13 H24 Cl2 N3 O4 P

CRN 88-12-0 CMF C6 H9 N O

RN 53899-91-5 HCA

CN Ethanaminium, N,N,N-trimethyl-2-[(2-methyl-1-oxo-2-propenyl)oxy]-, chloride, polymer with 3-[(2-chloroethyl)[3-(2-chloroethyl)tetrahydro-2-oxido-2H-1,3,2-oxazaphosphorin-2-yl]amino]propyl 2-methyl-2-propenoate (9CI) (CA INDEX NAME)

CM 1

CRN 53804-60-7 CMF C14 H25 Cl2 N2 O4 P

$$\begin{array}{c|c} \text{C1CH}_2-\text{CH}_2 & \text{CH}_2 \\ & \text{O} & \text{CH}_2 \\ & \text{N} & \text{//} & \text{|} & \text{|} \\ & \text{P} & \text{N}-\text{(CH}_2)_3-\text{O}-\text{C}-\text{C}-\text{Me} \\ & \text{|} & \text{|} & \text{|} \\ & \text{O} & \text{CH}_2-\text{CH}_2\text{C1} & \text{O} \\ \end{array}$$

CM 2

CRN 5039-78-1 CMF C9 H18 N O2 . Cl

$$\begin{array}{c} \text{O} \quad \text{CH}_2 \\ || \quad || \\ \text{Me}_3\text{+N}-\text{CH}_2-\text{CH}_2-\text{O}-\text{C}-\text{C}-\text{Me} \end{array}$$

• c1 -

(prepn. of)

CC 35-3 (Synthetic High Polymers) Section cross-reference(s): 1, 28, 32 IT 53899-92-6 (pharmacol. activity of) IT53803-65-9P 53803-72-8P 53803-73-9P 53803-74-0P 53803-75-1P 53803-76-2P 53804-59-4P 53804-61-8P 53804-63-0P 53804-65-2P 53804-66-3P **53804-67-4P 53804-68-5P** 53829-65-5P 53899-91-5P 54427-62-2P 54427-63-3P 54427-64-4P 54427-65-5P 54427-66-6P 54427-67-7P 54427-68-8P 54427-69-9P

10/500,565 ACQUAH

2 CM

110-85-0 CRN C4 H10 N2 CMF

ΝН

C08G IC

INCL 260047000CZ

37 (Plastics Fabrication and Uses) CC

Section cross-reference(s): 25 9.016-87-9, 111-40-0 **107-15-3**, 1,2-Ethanediamine 24802-03-7 Isocyanic acid, polymethylenepolyphenylene ester (microcapsules from polymers contg., manuf. of, by interfacial IT polymn.)

26680-16-0P 26680-17-1P IT

(microcapsules of, manuf. of, by interfacial polymn.)

ANSWER 19 OF 19 HCA COPYRIGHT 2006 ACS on STN 72:79882 Encapsulation with polymer material. Santo, John E.; Vandegaer, Jan E. (Pennwalt Corp.). U.S. US 3492380 APPLICATION: US (English). CODEN: USXXAM. 19700127, 3 pp.

Solid or liq. particles are encapsulated by an interfacial polycondensation reaction with a P-contg. polymer shell, which is acid resistant, but dissolves in bases. Thus, 5 g AΒ p-(chlorocarbonyl)phenyl reaction with a P-contg. polymer shell, which is acid resistant, but dissolves in bases. p-(chlorocarbonyl)phenyl dichlorophosphate (I) (prepd. by heating p-hydroxybenzoic acid with PCl5) in 5 ml C6H6 was added to an agitated soln. of 3.1 g hexamethylenediamine in 200 ml H2O to form solid shells, which ruptured under vacuum and released .apprx.50ml H2O. The shells were stable for 15 min in aq. solns. of pH <7, burst after 12 min at pH 7, and burst after 2 min at pH>7. A mixt. of I, polymethylene poly(phenylisocyanate) (PAPI), sebacoyl chloride, poly(vinyl alc.) and Thimet was added to a soln. of ethylenediamine, diethylenetriamine, and Na2CO3 to form an encapsulated insecticide. 2,2-Bis(4-hydroxyphenyl)propane bis-(dichlorophosphate), prepd. by adding PCl3 to a mixt. of bisphenol A and POCl3, was similarly used to encapsulate Diazinon. The use of p-(chlorocarbonylmethylene)phenyl dichlorophosphate instead of I is claimed.

IT 26680-16-0 26680-17-1

(Diazinon encapsulation with)

RN 26680-16-0 HCA

CN Phosphorodichloridic acid, 4-(chlorocarbonyl)phenyl ester, polymer with 1,6-hexanediamine (9CI) (CA INDEX NAME)

CM 1

CRN 20070-01-3 CMF C7 H4 Cl3 O3 P

CM 2

CRN 124-09-4 CMF C6 H16 N2

 $H_2N-(CH_2)_6-NH_2$

RN 26680-17-1 HCA

CN Phosphorodichloridic acid, 4-(chlorocarbonyl)phenyl ester, polymer with piperazine (9CI) (CA INDEX NAME)

CM 1

CRN 20070-01-3 CMF C7 H4 Cl3 O3 P

CRN 110-85-0 CMF C4 H10 N2

NH HN

IT 107-15-3, Ethylenediamine

(polymers with (chloroformyl) phenyl phosphorodichloridate and isocyanic acid esters, Thimet encapsulation with)

RN 107-15-3 HCA

CN 1,2-Ethanediamine (9CI) (CA INDEX NAME)

 $H_2N-CH_2-CH_2-NH_2$

IC A61J; B29C

INCL 264004000

CC 36 (Plastics Manufacture and Processing)

IT 111-19-3D, Sebacoyl chloride, polymers with diethylenetriamine and isocyanic acid esters 26680-16-0 26680-17-1

(Diazinon encapsulation with)

IT 107-15-3, Ethylenediamine 111-40-0 (polymers with (chloroformyl)phenyl phosphorodichloridate and isocyanic acid esters, Thimet encapsulation with)

=> d 196 1-47 ti

L96 ANSWER 1 OF 47 HCA COPYRIGHT 2006 ACS on STN

TI Poly(arylazophosphonate)s: new arylazophosphonate-containing monomers for synthesis of laser-structurable polymers

Poly(oxy-1,2-ethanediyl), .alpha.-(ethoxyphosphinyl)-.omega.-[(ethoxyphosphinyl)oxy] - (9CI) (CA INDEX NAME)

$$\begin{array}{c|c}
O & O & O \\
\parallel & O - CH_2 - CH_2 - O - PH - OEt
\end{array}$$

107-15-3, reactions IT

(with diethyl phosphite and paraformaldehyde)

107-15-3 HCA

1,2-Ethanediamine (9CI) (CA INDEX NAME) RNCN

 $_{\mathrm{H_2N^-CH_2^-CH_2^-NH_2}}$

C07F; C08G

36-6 (Plastics Manufacture and Processing) IC

40550-08-1 **39297-31-9** 40550-06-9 40550-07-0 CC IT

40623-48-1

(reaction with acetone and triethylenetetramine)

107-15-3, reactions IT

(with diethyl phosphite and paraformaldehyde)

ANSWER 18 OF 19 HCA COPYRIGHT 2006 ACS on STN 76:114359 Phosphorus-containing polymers. Santo, John E.; Vandegaer, Jan E. (Pennwalt Corp.). U.S. US 3624248 19711130, 4 pp. Division of U.S. 3,492,380 (CA 72;79882g). (English). CODEN: USXXAM. APPLICATION: US 1970-403 19700106.

A P-contg. solid polymer which is sol. in alk. but not acid media is prepd. by interfacial polycondensation of p-(chlorocarbonyl)phenyl dichlorophosphate (I) [20070-01-3] and diamines or triamines and AB used to encapsulate liq., solid, or gas, e.g. insecticides. 3.1 g hexamethylenediamine in 200 ml water is agitated with a soln. of 5 g I in 5 ml benzene for a few min and the capsules are sepd. The p-(chlorocarbonyl)phenyl dichlorophosphate-hexamethylenediamine polymer [26680-16-0] encapsulates .sim.50 ml water. The polymer shells rupture in vacuum, are stable for >15 min at pH .leq.4 but burst in 2 min at pH 8.

107-15-3, 1,2-Ethanediamine TI

(microcapsules from polymers contg., manuf. of, by interfacial polymn.)

107-15-3 HCA RN

1,2-Ethanediamine (9CI) (CA INDEX NAME) CN

 $_{\mathrm{H_2N^-CH_2^-CH_2^-NH_2}}$

ACQUAH 10/500,565

26680-16-0P 26680-17-1P

(microcapsules of, manuf. of, by interfacial polymn.)

26680-16-0 HCA RN

Phosphorodichloridic acid, 4-(chlorocarbonyl)phenyl ester, polymer CNwith 1,6-hexanediamine (9CI) (CA INDEX NAME)

1 CM

20070-01-3 CRN C7 H4 Cl3 O3 P CMF

2 CM

124-09-4 CRN C6 H16 N2 **CMF**

 H_2N^- (CH₂)₆-NH₂

26680-17-1 HCA RN

Phosphorodichloridic acid, 4-(chlorocarbonyl)phenyl ester, polymer CN with piperazine (9CI) (CA INDEX NAME)

CM 1

20070-01-3 CRN

C7 H4 Cl3 O3 P CMF